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Gerard CM, Baudson N, Kraemer K, Ledent C, Pardoll D, Bruck C.

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The early genes E6 and E7 of human papillomavirus type 16 (HPV16) are consistently and exclusively expressed in HPV16-induced cancer lesions and play major roles in the development and maintenance of the malignant phenotype. Because this protein is a good example of a tumor-associated antigen, we have used E7 as a model antigen to test the potential of an experimental vaccine as an immunotherapeutic approach. In this study, we used a murine E7-expressing tumor model (TC1 cells) to assess effects of an E7-based vaccine on tumor growth. We show that vaccination with the E7 protein, formulated in the SmithKline Beecham Biologicals proprietary adjuvants (SBAS 1 and SBAS 2), leads to the rejection of pre-established tumors. Tumor rejection was associated with the induction of a strong systemic T helper 1 response, including CTLs, and the presence of an inflammatory infiltrate within the regressing tumor. Because most identified tumor-associated antigens are self antigens rather viral antigens, we used E7 transgenic mice to evaluate the E7-based vaccine in conditions where E7 is a self antigen. Transgenic mice, which constitutively and specifically express the E7 HPV16 gene in the thyroid epithelium, rapidly develop thyroid goiters and, after several months, thyroid carcinomas. We show that E7-specific antibodies and CD4 T helper responses can be obtained by vaccinating E7 transgenic mice, although a CTL response was not detected. Despite the absence of measurable CTL responses, vaccination still reduced the growth of pre-established TC1 tumors, although less efficiently than in nontransgenic animals, but was unable to suppress or delay the development of the spontaneous thyroid pathology.

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, asmos	A DNA vaccine based on a shuffled E7 oncogene of the human papillomavirus type 16 (HPV 16) induces E7-specific cytotoxic T cells but lacks transforming activity.					
PubMed Services	Osen W, Peiler T, Ohlschlager P, Caldeira S, Faath S, Michel N, Muller M, Tommasino M, Jochmus I, Gissmann L.					
	Deutsches Krebsforschungszentrum, Angewandte Tumorvirologie Im Neuenheimer Feld 242, D-69120, Heidelberg, Germany.					
Related Resources	Vaccination with oncogene-derived DNA for anti-cancer treatment carries a risk of de-novo tumor induction triggered by the persisting recombinant DNA. We hypothesized that an oncoprotein whose primary sequence has been rearranged ('shuffled') to maintain all possible T cell epitopes still induces cytotoxic T cells against the authentic protein but is devoid of transforming properties. As a model antigen, we used the E7 oncoprotein of the human papillomavirus (HPV) type 16, the major cause of cervical cancer. We have generated an artificial E7 molecule in which four domains were rearranged and, in order to maintain all possible T cell epitopes, certain sequences were duplicated. Upon transfection of this shuffled E7 gene (E7SH) into RMA cells, presentation of an E7 Db-restricted T cell epitope was shown by an E7-specific CTL line in vitro. Immunization of C57BL/6 mice with E7SH DNA induced E7-specific CTL and also conveyed protection against E7-positive syngeneic tumor cells. No transforming activity of E7SH DNA in NIH3T3 cells was detected, as determined by focus formation, induction of S-phase under conditions of serum deprivation and degradation of endogenous pRB. Our results suggest that DNA shuffling may become a promising concept for DNA-based anti-cancer vaccines.					
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PubMed Services	tumour growth.  Gerard CM, Baudson N, Kraemer K, Bruck C, Garcon N, Paterson Y, Pan ZK, Pardoll D.				
30111003	SmithKline Beecham Biologicals, R&D, Preclinical Immunology, Rue de 1'Institut 89, B-1330, Rixensart, Belgium. catherine.gerard@sbbio.be  Over 90% of cervical cancers are associated with HPV infection, the commonest				
Related Resources	being the HPV-16 subtype. Two early viral genes, E6 and 7, play major roles in the development and maintenance of the malignant phenotype. The vaccine potential of a recombinant HPV16 E7 protein was examined in two murine models of E7-expressing tumours. Formulations including the immunostimulants MPL and QS21 induced therapeutically active immune responses leading to regression of pre-established TC1 tumour lesions, associated with induction of IgG antibodies, lymphoproliferation and CTL. Our data provide a clear incentive to investigate the clinical application of this approach in cancer immunotherapy.				
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PubMed Services	In vivo transfer of the human cystic fibrosis transmembrane conductance regulator gene to the airway epithelium.  Cell. 1992 Jan 10;68(1):143-55.  PMID: 1370653 [PubMed - indexed for MEDLINE]					
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        295493 "TUMOR"
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        335490 "TUMOR"
                 ("TUMOR" OR "TUMORS")
             0 "OR"
           948 "ORS"
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                 ("OR" OR "ORS")
        204831 "CANCER"
         29040 "CANCERS"
        212852 "CANCER"
                 ("CANCER" OR "CANCERS")
         41254 "VACCINE"
         42029 "VACCINES"
         51899 "VACCINE"
                 ("VACCINE" OR "VACCINES")
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L2
          1137 "TUMOR VACCINE"
                 ("TUMOR"(W) "VACCINE")
=> HPV and L2
          4768 HPV
           593 HPVS
          4806 HPV
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L3
            35 HPV AND L2
=> "early protein"
        348300 "EARLY"
            21 "EARLIES"
        348315 "EARLY"
                 ("EARLY" OR "EARLIES")
       1582536 "PROTEIN"
       1088253 "PROTEINS"
       1832740 "PROTEIN"
                 ("PROTEIN" OR "PROTEINS")
          1567 "EARLY PROTEIN"
L4
                 ("EARLY"(W) "PROTEIN")
=> L4 and L3
             1 L4 AND L3
L5
=> "E6 or E7"
          5012 "E6"
             0 "OR"
           948 "ORS"
           948 "OR"
                 ("OR" OR "ORS")
          4419 "E7"
L6
             0 "E6 OR E7"
                 ("E6"(W)"OR"(W)"E7")
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=> "E6" and L2
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            26 "E6" AND L2
L7
=> "E7" and L2
          4419 "E7"
            44 "E7" AND L2
=> L6 and L7
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L9
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         63848 "LIPOPROTEIN"
         70169 "LIPOPROTEINS"
         87045 "LIPOPROTEIN"
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       2057381 "D"
L10
           329 "LIPOPROTEIN D"
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=> L10 and L6
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=> "T helper epitope"
        706053 "T"
         22992 "HELPER"
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         23111 "HELPER"
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         32571 "EPITOPE"
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=> L13 and L7
            1 L13 AND L7
L14
=> L13 and L8
L15
            1 L13 AND L8
=> influenza and L7
         18018 INFLUENZA
            6 INFLUENZAS
         18020 INFLUENZA
                  (INFLUENZA OR INFLUENZAS)
L16
             1 INFLUENZA AND L7
=> infleunza and L8
             O INFLEUNZA
             0 INFLEUNZA AND L8
L17
=> influenza and L8
         18018 INFLUENZA
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                 (INFLUENZA OR INFLUENZAS)
L18
             2 INFLUENZA AND L8
=> CpG (w) motif
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7565 CPG 240 CPGS

7614 CPG

(CPG OR CPGS)

37704 MOTIF 65704 MOTIFS 89009 MOTIF

(MOTIF OR MOTIFS)

555 CPG (W) MOTIF

=> L19 and L7

L20 0 L19 AND L7

=> L19 and L8

0 L19 AND L8

=> L2 and L19

L22 6 L2 AND L19

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        312416 "HUMANS"
       1378875 "HUMAN"
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          6050 "PAPILLOMA"
          2105 "PAPILLOMAS"
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         63144 "VIRUSES"
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'E6' NOT FOUND
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=> envelope (w) protein
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          8488 ENVELOPES
         51077 ENVELOPE
                  (ENVELOPE OR ENVELOPES)
       1582536 PROTEIN
       1088253 PROTEINS
       1832740 PROTEIN
                  (PROTEIN OR PROTEINS)
L3
          9721 ENVELOPE (W) PROTEIN
=> L1 and 13
             9 L1 AND L3
=> CpG and L4
          7565 CPG
           240 CPGS
          7614 CPG
                  (CPG OR CPGS)
L5
             0 CPG AND L4
=> influenza and L1
         18018 INFLUENZA
             6 INFLUENZAS
         18020 INFLUENZA
                  (INFLUENZA OR INFLUENZAS)
L6
            19 INFLUENZA AND L1
=> HPV (w) antigen
          4768 HPV
           593 HPVS
```

4806 HPV

(HPV OR HPVS)

246813 ANTIGEN

196038 ANTIGENS

306186 ANTIGEN

(ANTIGEN OR ANTIGENS)

L7 39 HPV (W) ANTIGEN

=> influenza and L7

18018 INFLUENZA

6 INFLUENZAS

18020 INFLUENZA

(INFLUENZA OR INFLUENZAS)

L8 2 INFLUENZA AND L7

```
=> "T cell helper epitope"
        706053 "T"
       1730680 "CELL"
       1544002 "CELLS"
       2323964 "CELL"
                 ("CELL" OR "CELLS")
         22992 "HELPER"
           215 "HELPERS"
         23111 "HELPER"
                 ("HELPER" OR "HELPERS")
         32571 "EPITOPE"
         32336 "EPITOPES"
         48896 "EPITOPE"
                 ("EPITOPE" OR "EPITOPES")
L9
            16 "T CELL HELPER EPITOPE"
                 ("T"(W)"CELL"(W)"HELPER"(W)"EPITOPE")
=> L1 and L9
L10
             0 L1 AND L9
=> influenza and L9
         18018 INFLUENZA
             6 INFLUENZAS
         18020 INFLUENZA
                 (INFLUENZA OR INFLUENZAS)
             0 INFLUENZA AND L9
L11
=>
=>
=> "T helper epitope"
        706053 "T"
         22992 "HELPER"
           215 "HELPERS"
         23111 "HELPER"
                 ("HELPER" OR "HELPERS")
         32571 "EPITOPE"
         32336 "EPITOPES"
         48896 "EPITOPE"
                 ("EPITOPE" OR "EPITOPES")
L12
           196 "T HELPER EPITOPE"
                 ("T"(W)"HELPER"(W)"EPITOPE")
=> L12 and L1
             2 L12 AND L1
=> DIS L13 1- IBIB IABS
YOU HAVE REQUESTED DATA FROM 2 ANSWERS - CONTINUE? Y/(N):Y
THE ESTIMATED COST FOR THIS REQUEST IS 5.08 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y) /N:Y
L13 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN
                         2003:173012 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         138:203667
TITLE:
                         Long peptides of 22-40 amino acid residues that induce
                         and/or enhance antigen specific immune responses.
PATENT ASSIGNEE(S):
                         Leids Universitair Medisch Centrum, Neth.
SOURCE:
                         Eur. Pat. Appl., 50 pp.
                         CODEN: EPXXDW
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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=> "HPV E6 and/or E7"

L1 222 "HPV E6 AND/OR E7"

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L2 0 "PROTEIN D (L) INFLUENZA B"

=> "influenza B"

L3 2512 "INFLUENZA B"

=> L1 and L3

L4 0 L1 AND L3

=> CpG and L1

5 0 CPG AND L1

=> "immunostimulatory CpG oligonucleotide"

L6 26 "IMMUNOSTIMULATORY CPG OLIGONUCLEOTIDE"

=> L1 and L6

L7 0 L1 AND L6

=> "influenzae B"

L8 413 "INFLUENZAE B"

=> L1 and L8

L9 0 L1 AND L8

=> "helper epitope"

L10 392 "HELPER EPITOPE"

=> L8 and L10

L11 1 L8 AND L10

=> L1 and L11

L12 0 L1 AND L11

=> L1 and L10

L13 0 L1 AND L10

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                 MEDLINE Reload
         Jun 10
                  PCTFULL has been reloaded
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NEWS 12
                 USAN to be reloaded July 28, 2002;
NEWS 13
         Jul 22
                  saved answer sets no longer valid
NEWS 14
         Jul 29
                 Enhanced polymer searching in REGISTRY
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         Aug 08
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         Sep 16
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                  Indexing added to some pre-1967 records in CA/CAPLUS
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                  CA Section Thesaurus available in CAPLUS and CA
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L8 413 "INFLUENZAE B"

=> L1 and L8

L9 0 L1 AND L8

=> "helper epitope"

L10 392 "HELPER EPITOPE"

=> L8 and L10

L11 1 L8 AND L10

=> L1 and L11

L12 0 L1 AND L11

=> L1 and L10

L13 0 L1 AND L10

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=> L1 and L10

L13 0 L1 AND L10

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L9 0 L1 AND L8

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=> L8 and L10

L11 1 L8 AND L10

=> L1 and L11

L12 0 L1 AND L11

=> L1 and L10

L13 0 L1 AND L10

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L19
           210 LYTA
=> fusion partner
L20
          1512 FUSION PARTNER
=> L19 and L20
             2 L19 AND L20
L21
=> D L21 BIB TI SO AU ABS
    ANSWER 1 OF 2 CAPLUS COPYRIGHT 2001 ACS
L21
     1999:511245 CAPLUS
AN
     131:140508
DN
     Tumor-associated antigen derivatives of MAGE proteins and their use in
ΤI
     cancer vaccine therapy
     Cabezon, Silva Teresa; Cohen, Joseph; Slaoui, Moncef Mohamed; Vinals
IN
     Bassols, Carlota
PA
     Smithkline Beecham Biologicals S.A., Belg.; Cabezon Silva, Teresa
SO
     PCT Int. Appl., 74 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                           APPLICATION NO. DATE
PΙ
     WO 9940188
                      A2
                            19990812
                                           WO 1999-EP660
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             TJ, TM
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     GB 1998-2650
                      19980206
     WO 1999-EP660
                      19990202
TΙ
     Tumor-associated antigen derivatives of MAGE proteins and their use in
     cancer vaccine therapy
SO
     PCT Int. Appl., 74 pp.
     CODEN: PIXXD2
ΙN
     Cabezon, Silva Teresa; Cohen, Joseph; Slaoui, Moncef Mohamed; Vinals
     Bassols, Carlota
AΒ
     The present invention relates to derivs. of MAGE proteins and their use
in
     cancer vaccine therapy. In particular, the protein derivs. are: (1)
     fusion proteins comprising an antigen encoded by the MAGE family of
genes,
     linked to an immunol. fusion partner which provides T
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helper epitopes; (2) chem. modified MAGE proteins wherein the antigen's disulfide bridges are reduced and the resulting thiols blocked; and/or

(3) genetically modified MAGE proteins provided with an affinity tag and/or genetically modified to prevent disulfphide bridge formation. The preferred MAGE proteins are MAGE Al and MAGE A3. The fusion proteins of the invention comprise an immunol. fusion parter such as lipoprotein D from Haemophilus influenzae, the NS1 (hemagglutinin) non-structural protein from influenzae virus, and/or the Streptococcus pneumoniae protein

LYTA. In addn., novel methods are also described for purifying MAGE proteins and for formulating vaccines for treating a range of cancers. The fusion protein LPD-MAGE3-His was used, along with an adjuvant, in a vaccine for the treatment of melanoma, and a TH1 type immune response was raised against said compn. The novel MAGE protein purifn. process of the invention comprises reducing the disulfide bonds, blocking the resulting free thiol group with a blocking group, and subjecting the resulting deriv. to one or more chromatog. purifn. steps.

## => D L21 BIB TI SO AU ABS 2

fragment

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L21 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2001 ACS
    1999:468468 CAPLUS
ΑN
    131:86861
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TΙ
    E6 and E7 fusion proteins for vaccination against human papilloma virus
IN
    Dalemans, Wilfried L. J.; Gerard, Catherine Marie Ghislaine
PA
    Smithkline Beecham Biologicals S. A., Belg.
SO
    PCT Int. Appl., 62 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 2
    PATENT NO. KIND DATE APPLICATION NO. DATE
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    WO 9933868 A2 19990708
WO 9933868 A3 19990916
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                     A3 19990916
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A2 20001004 EP 1998-966706
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     EP 1040123
                                                           19981218
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     E6 and E7 fusion proteins for vaccination against human papilloma virus
TΙ
SO
     PCT Int. Appl., 62 pp.
     CODEN: PIXXD2
     Dalemans, Wilfried L. J.; Gerard, Catherine Marie Ghislaine
IN
AΒ
     The authors disclose the prepn. and characterization of fusion proteins
of
     E6 and/or E7 of human papilloma virus (type 16 or 18) linked to an
     immunol. fusion partner that provides Th1 cell-type
     help. In one example, using recombinant DNA technol., a fragment of
     protein D of Haemophilus influenzae B was fused to the N-terminal
```

of E6 and expressed in E. coli. In a second example, the immunol. **fusion partner** providing T-cell help is the **LytA** amidase of Streptococcus pneumoniae. Vaccination with a fusion protein, in combination with CpG oligonucleotide, induced the regression of HPV E6-mediated tumors.

=> log off

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ANSWER 1 OF 1 CAPLUS COPYRIGHT 2001 ACS
ΑN
     1997:97727 CAPLUS
DN
     Prophylactic and therapeutic vector vaccination using expression
TI
     constructs for individual epitopes of antigens
ΙN
     Weiner, David B.; Williams, William V.; Wang, Bin
     Wistar Institute, USA; Trustees of the University of Pennsylvania
PΑ
     U.S., 50 pp. Cont.-in-part of U.S. Ser. No. 29,336, abandoned.
SO
     CODEN: USXXAM
DT
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     WO 1994-US899
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     US 1995-495684
                      19950828
     Prophylactic and therapeutic vector vaccination using expression
ΤI
     constructs for individual epitopes of antigens
SO
     U.S., 50 pp. Cont.-in-part of U.S. Ser. No. 29,336, abandoned.
     CODEN: USXXAM
     Weiner, David B.; Williams, William V.; Wang, Bin
IN
    Methods of prophylactic and therapeutic immunization against infection,
AΒ
     hyperproliferative and autoimmune diseases are disclosed. An expression
     construct directing the synthesis of one or more epitopes, or analogs of
     epitopes, of an antigen is introduced into cells of an individual. The
     epitope is identical or substantially similar to an epitope of a pathogen
     antigen, a hyperproliferative cell assocd. protein or a protein assocd.
     with autoimmune disease resp. Methods of immunizing against HIV are
     described. Successful induction of immunity to HIV1 in mice by injection
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with an expression vector for the HIV-1 gene env.

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L10 ANSWER 28 OF 45 CAPLUS COPYRIGHT 2001 ACS
ΑN
    1986:532044 CAPLUS
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    105:132044
     Immunogenic complex and its use as an immune stimulant, vaccines
TI
    and reagent
ΙN
    Morein, Bror
PA
    Swed.
SO
    Eur. Pat. Appl., 65 pp.
    CODEN: EPXXDW
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    NO 168806
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PRAI SE 1984-5493 19841101
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EP 1986-906026 19861016 WO 1986-SE480 19861016 WO 1987-SE480 19870601

- TI Immunogenic complex and its use as an immune stimulant, vaccines and reagent
- and reagent
  SO Eur. Pat. Appl., 65 pp.
  CODEN: EPXXDW
- IN More

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L10 ANSWER 23 OF 45 CAPLUS COPYRIGHT 2001 ACS
AN
     1994:296664 CAPLUS
DN
     120:296664
ΤI
     Multiple immunogens in vaccines
ΙN
     Becker, Robert S.; Biscardi, Karen; Ferguson, Laura; Erdile, Lorne
PΑ
     Connaught Laboratories Inc., USA
SO
     Eur. Pat. Appl., 16 pp.
     CODEN: EPXXDW
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FAN.CNT 1
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ΤI
    Multiple immunogens in vaccines
SO
    Eur. Pat. Appl., 16 pp.
    CODEN: EPXXDW
ΙN
    Becker, Robert S.; Biscardi, Karen; Ferguson, Laura; Erdile, Lorne
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L10
    ANSWER 20 OF 45 CAPLUS COPYRIGHT 2001 ACS
     1995:742918 CAPLUS
ΑN
DN
     123:123177
ΤI
     Antigen-carbohydrate conjugates and their use in immunotherapy
     McKenzie, Ian Farquhar Campbell; Apostolopoulos, Vasso; Pietersz, Geoff
ΙN
     Allan
PΑ
     Austin Research Institute, Australia
     Eur. Pat. Appl., 34 pp.
SO
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     WO 1994-AU789
                      19941223
    Antigen-carbohydrate conjugates and their use in immunotherapy
TΙ
SO
     Eur. Pat. Appl., 34 pp.
     CODEN: EPXXDW
    McKenzie, Ian Farquhar Campbell; Apostolopoulos, Vasso; Pietersz, Geoff
IN
     Alla
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1

L10 ANSWER 17 OF 45 CAPLUS COPYRIGHT 2001 ACS AN 1996:369881 CAPLUS

DN 125:27699

TI Nucleic acids encoding mutant matrix proteins useful for attenuation or enhancement of influenza A virus

IN Kawaoka, Yoshihiro; Castrucci, Maria R.

PA St. Jude Children's Research Hospital, USA

SO PCT Int. Appl., 96 pp. CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

APPLICATION NO. DATE PATENT NO. KIND DATE FAIENI NO. KIND DATE -----ΡI WO 9610631 A1 19960411 WO 1995-US12357 19951002 W: AU, CA, JP RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE AU 9537278 A1 19960426 AU 1995-37278 19951002 PRAI US 1994-316419 19940930 US 1995-471100 19950606 WO 1995-US12357 19951002

TI Nucleic acids encoding mutant matrix proteins useful for attenuation or enhancement of influenza A virus

SO PCT Int. Appl., 96 pp. CODEN: PIXXD2

IN Kawa

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ANSWER 3 OF 45 CAPLUS COPYRIGHT 2001 ACS
     2000:155184 CAPLUS
DN
     132:204050
     Recombinant swinepox virus for expression of foreign antigens in
TI
     vaccine preparations
     Cochran, Mark D.; Junker, David E.
IN
PA
     Syntro Corporation, USA
     U.S., 262 pp., Cont.-in-part of U.S. Ser. No. 375,922.
     CODEN: USXXAM
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ΤI
     Recombinant swinepox virus for expression of foreign antigens in
     vaccine preparations
SO
     U.S., 262 pp., Cont.-in-part of U.S. Ser. No. 375,922.
     CODEN: USXXAM
ΙN
     Cochran, Mark D.; Junker, David E.
RE.CNT
       84
(1) Anon; EP 0284416 1988 CAPLUS
(2) Anon; WO 8903429 1989 CAPLUS
(5) Ben-Porat, T; Journal of Virology 1986, V154, P325 CAPLUS
(6) Bertholet, C; EMBO Journal 1986, V5, P1951 CAPLUS
(7) Bhat, R; Nucleic Acids Research 1989, V17, P1159 CAPLUS
ALL CITATIONS AVAILABLE IN THE RE FORMAT
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ANSWER 1 OF 1 CAPLUS COPYRIGHT 2001 ACS
     1997:97727 CAPLUS
DN
     126:156420
TI
     Prophylactic and therapeutic vector vaccination using expression
     constructs for individual epitopes of antigens
IN
     Weiner, David B.; Williams, William V.; Wang, Bin
     Wistar Institute, USA; Trustees of the University of Pennsylvania
PΑ
     U.S., 50 pp. Cont.-in-part of U.S. Ser. No. 29,336, abandoned.
SO
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     Prophylactic and therapeutic vector vaccination using expression
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     constructs for individual epitopes of antigens
     U.S., 50 pp. Cont.-in-part of U.S. Ser. No. 29,336, abandoned.
SO
     CODEN: USXXAM
    Weiner, David B.; Williams, William V.; Wang, Bin
ΙN
    Methods of prophylactic and therapeutic immunization against infection,
AB
    hyperproliferative and autoimmune diseases are disclosed. An expression
     construct directing the synthesis of one or more epitopes, or analogs of
     epitopes, of an antigen is introduced into cells of an individual. The
     epitope is identical or substantially similar to an epitope of a pathogen
    antigen, a hyperproliferative cell assocd. protein or a protein assocd.
    with autoimmune disease resp. Methods of immunizing against HIV are
    described. Successful induction of immunity to HIV1 in mice by injection
```

with an expression vector for the HIV-1 gene env.

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L3
    ANSWER 8 OF 16 CAPLUS COPYRIGHT 2001 ACS
ΑN
    1994:699108 CAPLUS
DN
    121:299108
ΤI
    Antigenic polypeptides from hemagglutinins conferring multistrain
immunity
     to influenza viruses A and B
IN
    Shatzman, Allan; Kane, James; Scott, Miller; Dillon, Susan
PA
    SmithKline Beecham Corp., USA
    PCT Int. Appl., 152 pp.
SO
    CODEN: PIXXD2
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PRAI US 1993-13415
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    US 1993-108914
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TΙ
    Antigenic polypeptides from hemagglutinins conferring multistrain
    to influenza viruses A and B
SO
    PCT Int. Appl., 152 pp.
    CODEN: PIXXD2
    Shatzman, Allan; Kane, James; Scott, Miller; Dillon, Susan
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IN Shatzman, Allan; Kane, James; Scott, Miller; Dillon, Susan Fusion proteins contg. sequences from the HA2 subunits of influenza virus hemagglutinins that are capable of inducing an immune response are described for use in vaccines. The preferred fusion partner in these proteins is another influenza virus protein, preferably NS1. The construction of a no. of fusion proteins and their manuf. by expression of the gene in Escherichia coli is described. Vaccines contg. three of these fusion proteins, with Al3+ and 3D-MPL as adjuvants were prepd. and used to inoculate mice at 0 and 21 days. At day 49, the mice were challenged with 3-5 LD50 of influenza virus. Mice inoculated with the mixed antigen showed 73-100% survival depending on the strain and mice inoculated with single antigens showed 0-80% survival with controls animals showing 0-7% survival.

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ΑN
     1999:166640 CAPLUS
DN
     130:222110
     Fusion proteins of human papillomavirus E6 and E7 stimulate a type 1
ΤI
     T-cell response
     Bruck, Claudine; Cabezon Silva, Teres; Delisse, Anne-Marie Eva Fernande;
IN
     Gerard, Catherine Marie Ghislaine; Lombardo-Bencheikh, Angela
     Smithkline Beecham Biologicals S.A., Belg.
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     PCT Int. Appl., 95 pp.
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ΤI
     Fusion proteins of human papillomavirus E6 and E7 stimulate a type 1
     T-cell response
SO
     PCT Int. Appl., 95 pp.
     CODEN: PIXXD2
     Bruck, Claudine; Cabezon Silva, Teres; Delisse, Anne-Marie Eva Fernande;
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     Gerard, Catherine Marie Ghislaine; Lombardo-Bencheikh, Angela
AΒ
     The authors disclose the plasmid construction, expression, and purifn.
     from E. coli of human papillomavirus early proteins E6 and E7 linked to
     immunol. active fusion partners. These fusion
     proteins elicit a Th1 helper cell response in immunized mice. Using an
     E6/E7 HPV-transformed epithelial cell line, a vaccine formulation
     prot
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ΑN
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TΙ
     Tumor-associated antigen derivatives of MAGE proteins and their use in
     cancer vaccine therapy
     Cabezon, Silva Teresa; Cohen, Joseph; Slaoui, Moncef Mohamed; Vinals
ΙN
     Bassols, Carlota
PΑ
     Smithkline Beecham Biologicals S.A., Belg.; Cabezon Silva, Teresa
     PCT Int. Appl., 74 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
     PATENT NO.
                                             APPLICATION NO.
                       KIND
                             DATE
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     WO 9940188
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ΤI
     Tumor-associated antigen derivatives of MAGE proteins and their use in
     cancer vaccine therapy
SO
     PCT Int. Appl., 74 pp.
     CODEN: PIXXD2
     Cabezon, Silva Teresa; Cohen, Joseph; Slaoui, Moncef Mohamed; Vinals
ΙN
     Bassols, Carlota
     The present invention relates to derivs. of MAGE proteins and their use
AΒ
in
     cancer vaccine therapy. In particular, the protein derivs. are: (1)
     fusion proteins comprising an antigen encoded by the MAGE family of
genes,
     linked to an immunol. fusion partner which provides T
     helper epitopes; (2) chem. modified MAGE proteins wherein the antigen's
     disulfide bridges are reduced and the the resulting thiols blocked;
     (3) genetically modified MAGE proteins provided with an affinity tag
     and/or genetically modified to prevent disulfphide bridge formation.
     preferred MAGE proteins are MAGE A1 and MAGE A3. The fusion proteins of
     the invention comprise an immunol. fusion parter such as
     {f lipoprotein}\ {f D} from Haemophilus influenzae, the
    NS1 (hemagglutinin) non-structural protein from influenzae virus,
     and/or the Streptococcus pneumoniae protein LYTA. In addn.,
     novel methods are also described for purifying MAGE proteins and for
     formulating vaccines for treating a range of cancers. The fusion protein
     LPD-MAGE3-His was used, along with an adjuvant, in a vaccine for the
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treatment of melanoma, and a TH1 type immune response was raised against said compn. The novel MAGE protein purifn. process of the invention comprises reducing the disulfide bonds, blocking the resulting free thiol group with a blocking group, and subjecting the resulting deriv. to one

or

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